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## UNITED STATES DEPARTMENT OF COMMERCE

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APPLICATION NO. FILING DATE FIRST NAMED INVENTOR ATTORNEY DOCKET NO. 09/301,507 04/28/99 CYNADER М 230018.40101 **EXAMINER** 000500 HM12/0921 SEED INTELLECTUAL PROPERTY LAW GROUP PLL MARTINELL, J 701 FIFTH AVE **ART UNIT** PAPER NUMBER **SUITE 6300** 

DATE MAILED:

1633

09/21/00

Please find below and/or attached an Office communication concerning this application or proceeding.

**Commissioner of Patents and Trademarks** 

## Application No. 09/301,507

Applicant(s)

Cynader et al

Examiner

Office Action Summary

**James Martinell** 

**Group Art Unit** 1633



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Responsive to communication(s) filed on	
☐ This action is <b>FINAL</b> .	
☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11; 453 O.G. 213.	
is longer, from the mailing date of this communication.	n is set to expire month(s), or thirty days, whichever Failure to respond within the period for response will cause the Extensions of time may be obtained under the provisions of
Disposition of Claims	
	is/are pending in the application.
Of the above, claim(s)	is/are withdrawn from consideration.
Claim(s)	is/are allowed.
☐ Claim(s)	is/are rejected.
	is/are objected to.
	are subject to restriction or election requirement.
Application Papers  See the attached Notice of Draftsperson's Patent The drawing(s) filed on	is approved disapproved.  iminer.  in priority under 35 U.S.C. § 119(a)-(d).  copies of the priority documents have been  derial Number)  from the International Bureau (PCT Rule 17.2(a)).
Attachment(s)  Notice of References Cited, PTO-892 Information Disclosure Statement(s), PTO-1449, Interview Summary, PTO-413 Notice of Draftsperson's Patent Drawing Review Notice of Informal Patent Application, PTO-152	
SEE OFFICE ACT	TION ON THE FOLLOWING PAGES

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Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I. Claim 1, drawn to a cDNA library from a kitten, classified in class 536, subclass 23.1.
- II. Claim 2, drawn to a cDNA library from a kitten and an adult cat, classified in class 536, subclass 23.1.
- III. Claim 3, drawn to a cDNA library from a dark reared adult cat, classified in class 536, subclass 23.1.
- IV. Claims 8-22, drawn to polynucleotides, classified in class 536, subclass 23.1.
- V. Claims 23-25, drawn to human genes that hybridize to certain SEQ ID NOs, classified in class 536, subclass 23.1.
- VI. Claim 26, drawn to antisense polynucleotides, classified in class 536, subclass 23.1.
- VII. Claim 27, drawn to triple helix probes, classified in class 536, subclass 23.1.
- VIII. Claims 40-54, drawn to peptides encoded by certain

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SEQ ID NOs, classified in class 424, subclasses 12 and 15.

IX. Claims 55 and 56, drawn to recombinant binding partners of various types, classified in class 530, subclass 387.1.

Claims 4-6 and 28-39 are ungrouped because they are improper multiple dependent claims.

The inventions are distinct, each from the other for the following reasons. The polynucleotides and cDNA libraries of Groups I-VII are materially different from and are therefore independent and distinct from the peptides and binding partners of Groups VIII and IX. The cDNA libraries of Groups I-III are independent and distinct from one another because they are from different sources and contain different cDNAs. Likewise, the polynucleotides identified by SEQ ID NOs in Group IV are independent and distinct from those of each of Groups I-III because Group IV contains different polynucleotides from any one of Groups I-III. The human genes of Group V are independent and

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distinct from the feline cDNA libraries of each of Groups I-III and the feline sequences of Group IV. The antisense polynucleotides of Group VI are independent and distinct from each of Groups I-V because the antisense polynucleotides are not required to have any sequences in common with the polynucleotides of Groups I-V. Likewise, the Group VII triple helix probes are independent and distinct from Groups I-VI because the triple helix probes are not required to have any sequences in common with the polynucleotides of Groups I-VI. The peptides of Group VIII are independent and distinct from the antibodies and recombinant binding partners of Group IX.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification and/or because of their recognized divergent subject matter restriction for examination purposes as indicated is proper.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be

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examined even though the requirement be traversed (37 CFR 1.143).

Additionally, should applicants elect any one of Groups IVIX, applicants are required to elect one nucleotide sequence or
one amino acid sequence as a reference sequence because each
nucleotide sequence is independent and distinct from every other
nucleotide sequence and each polypeptide sequence is independent
and distinct from every other polypeptide sequence.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Certain papers related to this application may be submitted to Group 1600 by facsimile transmission. Papers should be faxed to Art Unit 1633 at (703) 308-4242. The faxing of such papers must conform with the rules published in the Official Gazette,

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1156 OG 61 (November 16, 1993).

Any inquiry concerning this communication should be directed to J. Martinell at telephone number (703) 308-0296.

JAMES MARTINELL, Ph.D SENIOR LEVEL EXAMINER